Amendments to the Claims

WHAT IS CLAIMED IS:

1. (Original) A compound having a structural formula I,

or a pharmaceutically acceptable salt, solvate, hydrate or stereoisomer thereof, wherein: R^1 and R^2 are each independently: methyl or ethyl.

2. (Currently Amended) The compound of Claim 1, wherein the compound having a- is structural formula II,

$$O \longrightarrow \bigcup_{O \in \mathbb{N}^1} O \longrightarrow \bigcup_{O \in \mathbb{N}^2} O \cap \bigcup_{O \in$$

3. (Currently Amended) The compound of Claim 2, wherein the compound is (2S)-3-(4-{[2-(4-methoxy-phenyl)-ethylcarbamoyl]-methoxy}-phenyl)-2-methoxy-propionic acid of having a structural formula III,

4. (Currently Amended) The compound of Claim 1, wherein the compound is 3-(4-{[2-(4-ethoxy-phenyl)-ethylcarbamoyl]-methoxy}-phenyl)-2-methoxy-propionic acid of having a-structural formula IV,

5. (Currently Amended) The compound of Claim 4, wherein the compound is (S)-3-(4-{[2-(4-ethoxy-phenyl)-ethylcarbamoyl]-methoxy}-phenyl)-2-methoxy-propionic acid having a of structural formula V,

- 6. (Currently Amended) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a compound of Claims 1–5 Claim 1 or a pharmaceutically acceptable salt, solvate or hydrate thereof.
 - 7. (Currently Amended) A pharmaceutical composition comprising:
- (1) a compound of <u>Claims 1-5 Claim 1</u>, or a pharmaceutically acceptable salt, solvate, hydrate or stereoisomer thereof;
- (2) a second therapeutic agent selected from the group consisting of: insulin sensitizers, sulfonylureas, biguanides, meglitinides, thiazolidinediones, α-glucosidase inhibitors, insulin secretogogues, insulin, antihyperlipidemic agents, plasma HDL-raising agents, HMG-CoA reductase inhibitors, statins, acryl CoA:cholestrol acyltransferase inhibitors, antiobesity compounds, antihypercholesterolemic agents, fibrates, vitamins and aspirin; and
 - (3) optionally a pharmaceutically acceptable carrier.
- 8. (Withdrawn) A method of modulating a peroxisome proliferator activated receptor (PPAR) comprising the step of contacting the receptor with a compound of Claims 1-5, or a pharmaceutically acceptable salt, solvate or hydrate thereof.

- 9. (Canceled)
- 10. (Canceled)
- 11. (Canceled)
- 12. (Canceled)
- 13. (Canceled)
- 14. (Canceled)
- 15. (Canceled)
- 16. (Currently Amended) A method for lowering blood-glucose in a mammal comprising the step of administering an effective amount of a compound of Claims 1-5 Claim 1.
- 17. (Currently Amended) A method of treating disease or condition in a mammal selected from the group consisting of hyperglycemia, dyslipidemia, Type II diabetes, Type I diabetes, hypertriglyceridemia, syndrome X, insulin resistance, heart failure, diabetic dyslipidemia, hyperlipidemia, hypercholesteremia, hypertension, obesity, anorexia bulimia, anorexia nervosa, cardiovascular disease and other diseases where insulin resistance is a component, comprising the step of administering an effective amount of a compound of Claims 1-5 Claim 1.
- 18. (Currently Amended) A method of treating diabetes mellitus in a mammal comprising the step of administering to a mammal a therapeutically effective amount of a compound of <u>Claims 1-5 Claim 1</u>.
- 19. (Currently Amended) A method of treating cardiovascular disease in a mammal comprising the step of administering to a mammal a therapeutically effective amount of a compound of Claims 1–5 Claim 1, or a pharmaceutically acceptable salt, solvate or hydrate thereof.
- 20. (Currently Amended) A method of treating syndrome X in a mammal, comprising the step of administering to the mammal a therapeutically effective amount of a compound of Claims 1–5 Claim 1, or a pharmaceutically acceptable salt, solvate or hydrate thereof.

- 21. (Currently Amended) A method of treating a disease or condition in a mammal selected from the group consisting of hyperglycemia, dyslipidemia, Type II diabetes, Type I diabetes, hypertriglyceridemia, syndrome X, insulin resistance, heart failure, diabetic dyslipidemia, hyperlipidemia, hypercholesteremia, hypertension, obesity, anorexia bulimia, anorexia nervosa, cardiovascular disease and other diseases where insulin resistance is a component, comprising the step of administering an effective amount of a compound of Claims 1-5 Claim 1; and an effective amount of second therapeutic agent selected from the group consisting of: insulin sensitizers, sulfonylureas, biguanides, meglitinides, thiazolidinediones, α-glucosidase inhibitors, insulin secretogogues, insulin, antihyperlipidemic agents, plasma HDL-raising agents, HMG-CoA reductase inhibitors, statins, acryl CoA:cholestrol acyltransferase inhibitors, antiobesity compounds, antihypercholesterolemic agents, fibrates, vitamins and aspirin.
- 22. (Currently Amended) Use of a compound of <u>Claims 1-5 Claim 1</u>, or a pharmaceutically acceptable salt, solvate or hydrate thereof, for the manufacture of a medicament for the treatment of a condition modulated by a PPAR.
- 23. (Currently Amended) Use of a compound of <u>Claims 1-5 Claim 1</u>, or a pharmaceutically acceptable salt, solvate or hydrate thereof, for the manufacture of a medicament for the treatment of diabetes.